Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

1. (Original) A pharmaceutical granulated product having improved granulatability, which contains a pharmaceutical compound with poor wettability and a surfactant.

2. (Original) A granulated product, wherein the product contains a compound with poor wettability and a surfactant, and at least about 35% by weight with respect to the total weight of the product does not pass through a 100-mesh sieve.

3. (Original) The granulated product according to claim 2, wherein the weight ratio of the compound and the surfactant is 1 : about 0.001 to about 2.

4. (Original) The granulated product according to claim 3, wherein the weight ratio is 1 : about 0.001 to less than 1.

5. (Original) The granulated product according to claim 3, wherein the weight ratio is 1 : about 0.001 to less than 0.1.

6. (Original) The granulated product according to claim 3, wherein the weight ratio is 1 : about 0.005 to about 0.05.

7. (Original) The granulated product according to claim 2, wherein the ratio of the compound with respect to the total granulated product is about 20% by weight or more.

8. (Original) The granulated product according to claim 2, wherein the compound is a pharmaceutical compound.

9. (Currently amended) A molded product made by molding the granulated product according to ~~any one of claims 2 to 8~~ claim 2.

10. (Original) A method for improving granulatability of a pharmaceutical composition containing a pharmaceutical compound with poor wettability, which comprises adding a surfactant before or during the granulation.

11. (Original) A method for preparing a granulated product containing a compound with poor wettability, having improved granulatability, which comprises adding a surfactant

in the weight ratio of about 0.001 to about 2 with respect to the compound before or during the granulation.

12. (Original) The method according to claim 11, wherein a granulated product is obtained in which at least about 35% by weight with respect to the total weight of the product does not pass through a 100-mesh sieve.

13. (Original) The method according to claim 11, which involves wet granulation in a binder solution containing a surfactant.

14. (Original) The method according to claim 13, wherein the concentration of the surfactant in the binder solution is about 1 to about 1,000 mmol/L.

15. (Original) The method according to claim 13, wherein the concentration of the surfactant in the binder solution is about 10 to about 100 mmol/L.

16. (Original) The method according to claim 11, wherein the compound is a pharmaceutical compound.

17. (Currently amended) A method for preparing a molded product, comprising molding the granulated product which is obtained by the method according to ~~any one of claims 11 to 16~~ claim 11.

18. (Canceled)

19. (Canceled)